

Appendices

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Appendix A

Appendix A. Search Strategy

Ovid MEDLINE Search Strategy

- 1 Epidemiologic studies/ (5166)
- 2 exp case control studies/ (525207)
- 3 exp cohort studies/ (1131452)
- 4 Case control.tw. (57348)
- 5 (cohort adj (study or studies)).tw. (55643)
- 6 Cohort analy\$.tw. (2558)
- 7 (Follow up adj (study or studies)).tw. (32450)
- 8 (observational adj (study or studies)).tw. (27770)
- 9 Longitudinal.tw. (105249)
- 10 randomized controlled trial/ (316611)
- 11 clinical trial/ (468024)
- 12 clinical trial, phase i.pt. (11624)
- 13 clinical trial, phase ii.pt. (18360)
- 14 clinical trial, phase iii.pt. (6539)
- 15 clinical trial, phase iv.pt. (640)
- 16 controlled clinical trial.pt. (83472)
- 17 randomized controlled trial.pt. (316611)
- 18 multicenter study.pt. (136354)
- 19 clinical trial.pt. (468024)
- 20 or/1-19 (1821334)
- 21 Craniocerebral Trauma/ (17808)
- 22 exp Brain Injuries/ (42331)
- 23 Cerebrovascular Trauma/ (65)
- 24 brain injur*.ti,ab. (27162)
- 25 head injur*.ti,ab. (16984)
- 26 tbi.ti,ab. (9150)
- 27 or/21-26 (74809)
- 28 20 and 27 (13181)
- 29 Rehabilitation/ (15502)
- 30 rehab*.ti,ab. (86406)
- 31 neurorehabilitation.ti,ab. (736)
- 32 29 or 30 or 31 (92978)
- 33 28 and 32 (1350)
- 34 limit 33 to "all child (0 to 18 years)" (658)
- 35 limit 34 to "all adult (19 plus years)" (554)
- 36 33 not 34 (692)
- 37 35 or 36 (1246)
- 38 limit 37 to (addresses or autobiography or bibliography or biography or case reports or clinical conference or congresses or dictionary or directory or in vitro or interactive tutorial or interview or lectures or legal cases or legislation or news or newspaper article or patient education handout or periodical index or portraits or video-audio media or webcasts) (65)
- 39 37 not 38 (1181)
- 40 limit 39 to yr="1980 -Current" (1168)

PsycINFO Search Strategy

- 1 epidemiologic studies.mp. (8127)
- 2 case control.mp. (4559)
- 3 exp Longitudinal Studies/ (14968)
- 4 (cohort adj (study or studies)).tw. (6277)
- 5 Cohort analy\$.tw. (393)
- 6 (Follow up adj (study or studies)).tw. (8970)
- 7 (observational adj (study or studies)).tw. (3673)

- 8 longitudinal.mp. (60892)
- 9 randomized controlled trial.mp. (5151)
- 10 clinical trial.mp. or exp Clinical Trials/ (10383)
- 11 controlled clinical trial.mp. (745)
- 12 phase i clinical trial.mp. (17)
- 13 phase ii clinical trial.mp. (31)
- 14 phase iii clinical trial.mp. (32)
- 15 phase iv clinical trial.mp. (3)
- 16 multicenter study.mp. (710)
- 17 or/1-16 (103008)
- 18 exp Traumatic Brain Injury/ or exp Head Injuries/ or craniocerebral trauma.mp. (11296)
- 19 brain injur*.mp. (14000)
- 20 exp Cerebrovascular Accidents/ or cerebrovascular trauma.mp. (9891)
- 21 head injur*.mp. (5711)
- 22 tbi.mp. (4229)
- 23 or/18-22 (27560)
- 24 17 and 23 (1792)
- 25 exp Rehabilitation/ or exp Neuropsychological Rehabilitation/ or rehabilitation.mp. (67368)
- 26 rehab*.mp. (62993)
- 27 exp Neurorehabilitation/ or neurorehabilitation.mp. (588)
- 28 or/25-27 (69282)
- 29 24 and 28 (422)
- 30 limit 29 to (100 childhood <birth to age 12 yrs> or 120 neonatal <birth to age 1 mo> or 140 infancy <age 2 to 23 mo> or 160 preschool age <age 2 to 5 yrs> or 180 school age <age 6 to 12 yrs> or 200 adolescence <age 13 to 17 yrs>) (77)
- 31 limit 30 to ("300 adulthood <age 18 yrs and older>" or 320 young adulthood <age 18 to 29 yrs> or 340 thirties <age 30 to 39 yrs> or 360 middle age <age 40 to 64 yrs> or "380 aged <age 65 yrs and older>" or "390 very old <age 85 yrs and older>") (66)
- 32 29 not 30 (345)
- 33 31 or 32 (411)
- 34 limit 33 to yr="1980 -Current" (409)

Cochrane Central Register of Controlled Trials Search Strategy

- 1 traumatic brain injur* and rehab* (224)

PEDro Search Strategy

- 1 traumatic brain injur* AND rehab* (34)

Appendix B

Appendix B. Table 1. Risk of Bias for Individual Studies

Study	Study design	Overall Risk of Bias Assessment	Comments
Cicerone, 2008 ¹	RCT	Moderate	Possible contamination via same professionals delivering treatment and control interventions; minimally clinically important difference in CIQ not specified <i>a priori</i> ; subjective self-report scale used for primary outcome measurement; no adjustment for multiple comparisons.
Vanderploeg, 2008 ²	RCT	Low	Well-designed study; no adjustment for multiple comparisons.
Salazar, 2000 ³	RCT	Moderate	Outcome assessors not blinded; intervention implementation judged partially adequate; primary outcomes self-report; no adjustment for multiple comparisons.
Greenwood, 1994 ⁴	RCT	Moderate	Group randomization; moderate attrition at 6-month time point, high attrition at 12-month time point; no adjustment for multiple comparisons. Outcomes at 24 months considered high risk of bias due to high attrition and not used.
Ponsford, 2006 ⁵	Cohort	High	Potential selection bias, retrospective control group; intervention definition and implementation partially adequate; no adjustment for multiple comparisons, many outcomes assessed including several scales and subscales; potential reporting bias.
Sarajuuri, 2005 ⁶	Cohort	Moderate	Potential selection bias; confounding not appropriately addressed.
Prigatano, 1994 ⁷	Cohort	High	Potential selection bias, retrospective control group; outcome assessors not blinded; intervention implementation partially adequate; inconsistent outcomes measurement across groups; confounding not adequately addressed.
Rattok, 1992 ⁸	Cohort	Moderate	Possible contamination via same professionals delivering treatment and control interventions; blinding of outcomes assessors not reported; no adjustment for multiple comparisons.
Prigatano, 1984 ⁹	Cohort	Moderate	Potential selection bias, retrospective control group; inadequate intervention implementation; inconsistent outcomes measurement across groups; confounding not adequately addressed.
Hashimoto, 2006 ¹⁰	Cohort	High	Potential selection bias; blinding of outcomes assessors not reported, inadequate intervention definition; treatment group provided varying levels of treatment intensity, but comparisons are for entire group to a no treatment group; subjective self-report scale used for primary outcome measurement; minimally clinically important difference in CIQ not specified <i>a priori</i> ; confounding not adequately addressed; no adjustment for multiple comparisons, many outcomes assessed including several scales and subscales.
Cicerone, 2004 ¹¹	Cohort	Moderate	Selection bias; intervention definition and implementation partially adequate; subjective self-report scale used for primary outcome measurement; confounding not adequately addressed; no adjustment for multiple comparisons.
Willer, 1999 ¹²	Cohort	High	Potential selection bias; inadequate intervention definition; intervention implementation partially adequate; subjective self-report scale used for primary outcome measurement; minimally clinically important difference in CIQ not

Study	Study design	Overall Risk of Bias Assessment	Comments
			specified <i>a priori</i> ; insufficient statistical analysis; confounding not adequately addressed; no adjustment for multiple comparisons.
Bell, 2005 ¹³	RCT	NA	Studies with only secondary outcomes not assessed for risk of bias.
Powell, 2002 ¹⁴	RCT	NA	Studies with only secondary outcomes not assessed for risk of bias.
Thomas, 2004 ¹⁵	Cohort	NA	Studies with only secondary outcomes not assessed for risk of bias.
Semlyen, 1998 ¹⁶	Cohort	NA	Studies with only secondary outcomes not assessed for risk of bias.

Appendix B. Table 2. Risk of Bias Assessment Form for RCTs

Author _____ Year _____ PMID _____ Reviewer _____

Question	Response	Criteria	Justification
Internal Validity			
1. Was the method of randomization adequate?	Yes <input type="checkbox"/>	Method used should produce comparable groups.	
	No <input type="checkbox"/>	Pseudo randomization (ie. alternate allocation, by days of week, etc) or randomization approach cannot be determined	
	Uncertain <input type="checkbox"/>	Randomization method unclear	
2. Was allocation concealment adequate?	Yes <input type="checkbox"/>	Method used to conceal the allocation sequence could not have been foreseen in advance of, or during, enrolment.	
	No <input type="checkbox"/>	No concealment	
	Uncertain <input type="checkbox"/>	Could not be ascertained.	
3. Were outcome assessors blinded?	Yes <input type="checkbox"/>	Yes	
	No <input type="checkbox"/>	No	
	Uncertain <input type="checkbox"/>	Could not be ascertained.	
4a. Is the level of detail in describing the treatment intervention adequate?	Yes <input type="checkbox"/>	Treatment intervention described based upon model or theory, specific intervention components adequately described, interventions documented in manuals or other documentation.	
	Partially <input type="checkbox"/>	Some of the above features.	
	No <input type="checkbox"/>	None of the above features.	
4b. Is the level of detail in describing the control intervention adequate?	Yes <input type="checkbox"/>	Active control intervention described based upon model or theory, specific intervention components adequately described, interventions documented in manuals or other documentation. Passive control adequately described.	
	Partially <input type="checkbox"/>	Some of the above features.	
	No <input type="checkbox"/>	None of the above features.	
5. Are interventions assessed using valid and reliable measures, implemented consistently across all study participants?	Yes <input type="checkbox"/>	Implementation accompanied by staff training and fidelity checks, consistency across groups in treatment features not studied.	
	Partially <input type="checkbox"/>	Implementation accompanied by some of above features.	
	No <input type="checkbox"/>	No training or fidelity checks.	
6. Are outcomes assessed using valid and reliable measures, implemented	Yes <input type="checkbox"/>	Measure valid and reliable (i.e. objective measures, well validated scale, provider report)	
	Partially <input type="checkbox"/>	Some of the above features	

consistently across all study participants?		(partially validated scale)	
	No <input type="checkbox"/>	None of the above features. (self-report, scales with lower validity, reliability)	
7. Were incomplete outcome data adequately addressed?	Yes <input type="checkbox"/>	Balanced across groups and/or imputed using appropriate methods.	
	No <input type="checkbox"/>	High attrition or differential loss; no imputations or inappropriate imputations for missing data.	
	Uncertain <input type="checkbox"/>	Could not be ascertained.	
8. Are reports of the study free of suggestion of selective outcome reporting?	Yes <input type="checkbox"/>	All prespecified outcomes reported.	
	No <input type="checkbox"/>	Not all prespecified outcomes reported, subscales reported not prespecified, outcomes reported incompletely.	
	Uncertain <input type="checkbox"/>	Could not be ascertained.	
9. Is the study free from additional sources of bias?	Yes <input type="checkbox"/>		
	No <input type="checkbox"/>		
	Uncertain <input type="checkbox"/>		
	Overall Assessment		
Overall Risk of Bias assessment	Low <input type="checkbox"/>	Results are believable taking study limitations into consideration	
	Moderate <input type="checkbox"/>	Results are probably believable taking study limitations into consideration	
	High <input type="checkbox"/>	Results are uncertain taking study limitations into consideration	

Appendix B. Table 3. Risk of Bias Assessment Form for Observational Studies

Author _____ Year _____ PMID _____ Reviewer _____

Question	Response	Criteria	Justification
Internal Validity			
1. Is the study design prospective, retrospective, or mixed?	Prospective <input type="checkbox"/>	Outcome has not occurred at the time the study is initiated and information is collected over time to assess relationships with the outcome.	
	Mixed <input type="checkbox"/>	Case-control or cohort studies in which one group is studied prospectively and the other retrospectively.	
	Retrospective <input type="checkbox"/>	Analyzes data from past records.	
2a. Are inclusion/exclusion criteria clearly stated (i.e., severity, time since injury, pre-existing conditions, comorbidities, prior tbi)	Yes <input type="checkbox"/>		
	Partially <input type="checkbox"/>	Some, but not all, criteria stated or some not clearly stated.	
	No <input type="checkbox"/>		
2b. TBI severity inclusion criteria measured using valid and reliable measures and appropriate cut points for mod/sev TBI?	Yes <input type="checkbox"/>	e.g., GCS<13; LOC> 30 minutes; AOC >24 hours; PTA>1 day; AISS>2; positive imaging	
	No <input type="checkbox"/>		
	Uncertain <input type="checkbox"/>	Could not be ascertained.	
2c. Did the study apply inclusion/exclusion criteria uniformly to all comparison groups of the study?	Yes <input type="checkbox"/>		
	Partially <input type="checkbox"/>	Some criteria applied to all arms	
	No <input type="checkbox"/>		
2d. Is the selection of the comparison group appropriate, after taking into consideration feasibility and ethical considerations?	Yes <input type="checkbox"/>	Groups selected from same source (e.g., community or hospital) to reduce baseline differences between groups. For case-control studies, cases should have met case definition if they had the outcome.	
	No <input type="checkbox"/>		
	Uncertain <input type="checkbox"/>	Could not be ascertained.	
3. Were outcome assessors blinded?	Yes <input type="checkbox"/>	Yes	
	No <input type="checkbox"/>	No	
	Uncertain <input type="checkbox"/>	Could not be ascertained.	
4a. Is the level of detail in describing the treatment intervention adequate?	Yes <input type="checkbox"/>	Treatment intervention described based upon model or theory, specific intervention components adequately described, interventions documented in manuals or other documentation.	
	Partially <input type="checkbox"/>	Some of the above features.	

	No <input type="checkbox"/>	None of the above features.	

4b. Is the level of detail in describing the control intervention adequate?	Yes <input type="checkbox"/>	Intervention described based upon model or theory, specific intervention components adequately described, interventions documented in manuals or other documentation.	
	Partially <input type="checkbox"/>	Some of the above features.	
	No <input type="checkbox"/>	None of the above features.	
5. Are interventions assessed using valid and reliable measures, implemented consistently across all study participants?	Yes <input type="checkbox"/>	Implementation accompanied by staff training and supervision, checks of adherence/fidelity; consistency across groups in treatment features not studied.	
	Partially <input type="checkbox"/>	Implementation accompanied by some of above features.	
	No <input type="checkbox"/>	Implementation accompanied by none of above features.	
6. Are outcomes assessed using valid and reliable measures, implemented consistently across all study participants?	Yes <input type="checkbox"/>	Measure valid and reliable (i.e. objective measures, well validated scale, provider report); consistent implementation across groups.	
	Partially <input type="checkbox"/>	Some of the above features (partially validated scale)	
	No <input type="checkbox"/>	None of the above features. (self-report, scales with lower validity, reliability); in consistent implementation across groups	
	Uncertain <input type="checkbox"/>	Could not be ascertained.	
7a. Was attrition from all groups less than 20 percent?	Yes <input type="checkbox"/>		
	No <input type="checkbox"/>		
	Uncertain <input type="checkbox"/>	Could not be ascertained (i.e. retrospective designs where eligible at baseline could not be determined)	
7b. Did attrition differ between groups by less than 20 percent?	Yes <input type="checkbox"/>		
	No <input type="checkbox"/>		
	Uncertain <input type="checkbox"/>	Could not be ascertained (i.e. retrospective designs where eligible at baseline could not be determined)	
7c. In cases of high attrition or differential attrition, is the impact assessed (e.g. through sensitivity analysis or other adjustment method)?	Yes <input type="checkbox"/>		
	No <input type="checkbox"/>		
	Uncertain <input type="checkbox"/>	Could not be ascertained (i.e. retrospective designs where eligible at baseline could not be determined)	

	NA <input type="checkbox"/>	Not considered high or case-control study	
8. Were the important confounding and effect modifying variables taken into account in the design and/or analysis (e.g. through matching, stratification, interaction terms, multivariate analysis, or other statistical adjustment)?	Yes <input type="checkbox"/>		
	Partially <input type="checkbox"/>	Some variables taken into account or adjustment achieved to some extent	
	No <input type="checkbox"/>	Not accounted for or not identified.	
	Uncertain <input type="checkbox"/>	Could not be ascertained	
9. Are the statistical methods used to assess the primary outcomes appropriate to the data?	Yes <input type="checkbox"/>	Statistical techniques used must be appropriate to the data and take into account issues such as controlling for dose-response, small sample size, clustering, rare outcomes, and multiple comparisons. In normally distributed data the standard error, standard deviation, or confidence intervals should be reported. In non-normally distributed data, inter-quartile range should be reported.	
	Partially <input type="checkbox"/>		
	No <input type="checkbox"/>		
	Uncertain <input type="checkbox"/>	Could not be ascertained	
10. Are reports of the study free of suggestion of selective outcome reporting?	Yes <input type="checkbox"/>		
	No <input type="checkbox"/>	Not all prespecified outcomes reported, subscales not prespecified reported, outcomes reported incompletely.	
	Uncertain <input type="checkbox"/>	Could not be ascertained.	
11. Is the study free from additional sources of bias?	Yes <input type="checkbox"/>		
	No <input type="checkbox"/>		
	Uncertain <input type="checkbox"/>		
	Overall Assessment		
Overall Risk of Bias assessment	Low <input type="checkbox"/>	Results are believable taking study limitations into consideration	
	Moderate <input type="checkbox"/>	Results are probably believable taking study limitations into consideration	
	High <input type="checkbox"/>	Results are uncertain taking study limitations into consideration	

Appendix C

Appendix C. Studies undergoing full text review

1. Altman IM, Swick S, Parrot D, et al. Effectiveness of community-based rehabilitation after traumatic brain injury for 489 program completers compared with those precipitously discharged. *Archives of Physical Medicine & Rehabilitation*. 2010 Nov;91(11):1697-704. 21044714. *Not eligible study design*
2. Anderson SI, Wilson CL, McDowell IP, et al. Late rehabilitation for closed head injury: a follow-up study of patients 1 year from time of discharge. *Brain Injury*. 1996 Feb;10(2):115-24. 8696311. *No comparison group*
3. Ashley MJ, Persel CS, Clark MC, et al. Long-term follow-up of post-acute traumatic brain injury rehabilitation: a statistical analysis to test for stability and predictability of outcome. *Brain Injury*. 1997 Sep;11(9):677-90. 9376835. *Not intervention study*
4. Ashley MJ, Persel CS, Lehr RP, Jr., et al. Post-acute rehabilitation outcome: relationship to case-management techniques and strategy. *Journal of Insurance Medicine (Seattle)*. 1994;26(3):348-54. 10150511. *Not eligible study design*
5. Backhaus SL, Ibarra SL, Klyce D, et al. Brain injury coping skills group: a preventative intervention for patients with brain injury and their caregivers.[Erratum appears in Arch Phys Med Rehabil. 2010 Nov;91(11):1793]. *Archives of Physical Medicine & Rehabilitation*. 2010 Jun;91(6):840-8. 20510972. *No primary or secondary outcomes*
6. Bateman A, Culpan FJ, Pickering AD, et al. The effect of aerobic training on rehabilitation outcomes after recent severe brain injury: a randomized controlled evaluation. *Archives of Physical Medicine & Rehabilitation*. 2001 Feb;82(2):174-82. 11239307. *No primary or secondary outcomes*
7. Bell KR, Temkin NR, Esselman PC, et al. The effect of a scheduled telephone intervention on outcome after moderate to severe traumatic brain injury: a randomized trial. *Archives of Physical Medicine & Rehabilitation*. 2005 May;86(5):851-6. 15895327. *Eligible*
8. Benge JF, Caroselli JS, Reed K, et al. Changes in supervision needs following participation in a residential post-acute brain injury rehabilitation programme. *Brain Injury*. 2010;24(6):844-50. 20377342. *Not eligible comparison group*
9. Bornhofen C, McDonald S. Comparing strategies for treating emotion perception deficits in traumatic brain injury. *Journal of Head Trauma Rehabilitation*. 2008 Mar-Apr;23(2):103-15. 18362764. *Impairment-specific intervention*
10. Bornhofen C, McDonald S. Treating deficits in emotion perception following traumatic brain injury. *Neuropsychological Rehabilitation*. 2008 Jan;18(1):22-44. 17852760. *Impairment-specific intervention*
11. Bourgeois MS, Lenius K, Turkstra L, et al. The effects of cognitive teletherapy on reported everyday memory behaviours of persons with chronic traumatic brain injury. *Brain Injury*. 2007 Nov;21(12):1245-57. 18236200. *Not 75% moderate/severe TBI*
12. Bowen A, Tennant A, Neumann V, et al. Neuropsychological rehabilitation for traumatic brain injury: do carers benefit? *Brain Injury*. 2001 Jan;15(1):29-38. 11201312. *No primary or secondary outcomes*
13. Braunling-McMorrow D, Dollinger SJ, Gould M, et al. Outcomes of post-acute rehabilitation for persons with brain injury. *Brain Injury*. 2010;24(7-8):928-38. 20545448. *No comparison group*
14. Braverman SE, Spector J, Warden DL, et al. A multidisciplinary TBI inpatient rehabilitation programme for active duty service members as part of a randomized clinical trial. *Brain Injury*. 1999 Jun;13(6):405-15. 10401542. *Eligible - companion to 10865301*
15. Brooks N. The effectiveness of post-acute rehabilitation. *Brain Injury*. 1991 Apr-Jun;5(2):103-9. 1873599. *No original data*
16. Burke WH, Wesolowski MD, Guth ML. Comprehensive head injury rehabilitation: an outcome evaluation. *Brain Injury*. 1988 Oct-Dec;2(4):313-22. 3203177. *No comparison group*
17. Bush BA, Novack TA, Malec JF, et al. Validation of a model for evaluating outcome after traumatic brain injury. *Archives of Physical Medicine & Rehabilitation*. 2003 Dec;84(12):1803-7. 14669187. *No comparison group*
18. Cannon XL, Zhu WS, Poon Chetwyn CCCSW. Does Intensive Rehabilitation Improve Functional Outcome In Patients with Traumatic Brain Injury (TBI). Preliminary Results of a Prospective Randomized Controlled Trial. *Journal of Neurotrauma*. 1998(1):85. CN-00689851. *No primary or secondary outcomes*
19. Carnevale GJ, Anselmi V, Busichio K, et al. Changes in ratings of caregiver burden following a community-based behavior management program for persons with traumatic brain injury. *The Journal of head trauma rehabilitation*. 2002(2):83-95. CN-00378995. *Not 75% moderate/severe TBI*

20. Carnevale GJ, Anselmi V, Johnston MV, et al. A natural setting behavior management program for persons with acquired brain injury: a randomized controlled trial. *Archives of physical medicine and rehabilitation*. 2006(10):1289-97. CN-00568342. *No primary or secondary outcomes*
21. Cattelani R, Roberti R, Lombardi F. Adverse effects of apathy and neurobehavioral deficits on the community integration of traumatic brain injury subjects. *European journal of physical & rehabilitation medicine*. 2008 Sep;44(3):245-51. 18762734. *Not intervention study*
22. Cattelani R, Tanzi F, Lombardi F, et al. Competitive re-employment after severe traumatic brain injury: clinical, cognitive and behavioural predictive variables. *Brain Injury*. 2002 Jan;16(1):51-64. 11796099. *Not intervention study*
23. Cattelani R, Zettin M, Zoccolotti P. Rehabilitation treatments for adults with behavioral and psychosocial disorders following acquired brain injury: a systematic review. *Neuropsychology Review*. 2010 Mar;20(1):52-85. 20143264. *No original data*
24. Chang Zj LP. Rehabilitation and acupuncture treatment for patients with traumatic brain injury. *Chinese Journal of Medical Device*. 2005(5):38-9. CN-00784100. *No primary or secondary outcomes*
25. Chard SE. Community neurorehabilitation: a synthesis of current evidence and future research directions. *NeuroRx*. 2006 Oct;3(4):525-34. 17012066. *No original data*
26. Chen SH, Thomas JD, Glueckauf RL, et al. The effectiveness of computer-assisted cognitive rehabilitation for persons with traumatic brain injury. *Brain Injury*. 1997 Mar;11(3):197-209. 9058001. *No primary or secondary outcomes*
27. Chesnut RM, Carney N, Maynard H, et al. Summary report: evidence for the effectiveness of rehabilitation for persons with traumatic brain injury. *The Journal of Head Trauma Rehabilitation* 1999;14(2):176-188. 1999. *No original data*
28. Choi JH, Jakob M, Stapf C, et al. Multimodal early rehabilitation and predictors of outcome in survivors of severe traumatic brain injury. *Journal of Trauma-Injury Infection & Critical Care*. 2008 Nov;65(5):1028-35. 19001970. *No comparison group*
29. Cicerone KD, Azulay J, Trott C. Methodological quality of research on cognitive rehabilitation after traumatic brain injury. *Archives of Physical Medicine & Rehabilitation*. 2009 Nov;90(11 Suppl):S52-9. 19892075. *No original data*
30. Cicerone KD, Dahlberg C, Kalmar K, et al. Evidence-based cognitive rehabilitation: recommendations for clinical practice. *Archives of Physical Medicine & Rehabilitation*. 2000 Dec;81(12):1596-615. 11128897. *No original data*
31. Cicerone KD, Dahlberg C, Malec JF, et al. Evidence-based cognitive rehabilitation: updated review of the literature from 1998 through 2002. *Archives of Physical Medicine & Rehabilitation*. 2005 Aug;86(8):1681-92. 16084827. *No original data*
32. Cicerone KD, Langenbahn DM, Braden C, et al. Evidence-based cognitive rehabilitation: updated review of the literature from 2003 through 2008. *Archives of Physical Medicine & Rehabilitation*. 2011 Apr;92(4):519-30. 21440699. *No original data*
33. Cicerone KD, Mott T, Azulay J, et al. Community integration and satisfaction with functioning after intensive cognitive rehabilitation for traumatic brain injury. *Archives of Physical Medicine & Rehabilitation*. 2004 Jun;85(6):943-50. 15179648. *Eligible*
34. Cicerone KD, Mott T, Azulay J, et al. A randomized controlled trial of holistic neuropsychologic rehabilitation after traumatic brain injury. *Archives of Physical Medicine & Rehabilitation*. 2008 Dec;89(12):2239-49. 19061735. *Eligible*
35. Cifu DX, Kreutzer JS, Kolakowsky-Hayner SA, et al. The relationship between therapy intensity and rehabilitative outcomes after traumatic brain injury: a multicenter analysis. *Archives of Physical Medicine & Rehabilitation*. 2003 Oct;84(10):1441-8. 14586910. *No primary or secondary outcomes*
36. Coetzer R, Rushe R. Post-acute rehabilitation following traumatic brain injury: are both early and later improved outcomes possible? *International Journal of Rehabilitation Research*. 2005 Dec;28(4):361-3. 16319563. *No comparison group*
37. Constantinidou F, Thomas RD, Robinson L. Benefits of categorization training in patients with traumatic brain injury during post-acute rehabilitation: additional evidence from a randomized controlled trial. *Journal of Head Trauma Rehabilitation*. 2008 Sep-Oct;23(5):312-28. 18815508. *Impairment-specific intervention*

38. Cope DN, Cole JR, Hall KM, et al. Brain injury: analysis of outcome in a post-acute rehabilitation system. Part 2: Subanalyses. *Brain Injury*. 1991 Apr-Jun;5(2):127-39. 1908341. *No comparison group*
39. Cope DN, Cole JR, Hall KM, et al. Brain injury: analysis of outcome in a post-acute rehabilitation system. Part 1: General analysis. *Brain Injury*. 1991 Apr-Jun;5(2):111-25. 1873600. *No comparison group*
40. Cusick CP, Gerhart KA, Mellick D, et al. Evaluation of the home and community-based services brain injury Medicaid Waiver Programme in Colorado. *Brain Injury*. 2003 Nov;17(11):931-45. 14514446. *Not eligible study design*
41. Dahlberg CA, Cusick CP, Hawley LA, et al. Treatment efficacy of social communication skills training after traumatic brain injury: a randomized treatment and deferred treatment controlled trial. *Archives of Physical Medicine & Rehabilitation*. 2007 Dec;88(12):1561-73. 18047870. *Impairment-specific intervention*
42. Dawson DR. A multidisciplinary community-based rehabilitation program improved social functioning in severe traumatic brain injury. *ACP Journal Club*. 2002(1):22. CN-00477567. *No original data*
43. Devitt R, Colantonio A, Dawson D, et al. Prediction of long-term occupational performance outcomes for adults after moderate to severe traumatic brain injury. *Disability & Rehabilitation*. 2006 May 15;28(9):547-59. 16690584. *Not intervention study*
44. Dirette DK, Hinojosa J, Carnevale GJ. Comparison of remedial and compensatory interventions for adults with acquired brain injuries. *Journal of Head Trauma Rehabilitation*. 1999 Dec;14(6):595-601. 10671705. *Not 75% moderate/severe TBI*
45. Do HK, Sahagian DA, Schuster LC, et al. Head trauma rehabilitation: program evaluation. *Rehabilitation Nursing*. 1988 Mar-Apr;13(2):71-5. 3353569. *No primary or secondary outcome*
46. Doig E, Fleming J, Tooth L. Patterns of community integration 2-5 years post-discharge from brain injury rehabilitation. *Brain Injury*. 2001 Sep;15(9):747-62. 11516344. *Not intervention study*
47. Drechsler R, Padovan F, Di Stefano G, et al. [An integrated concept for vocational rehabilitation of brain injured patients--a catamnestic study of occupational outcome 1 to 2 years later]. *Rehabilitation*. 1995 Nov;34(4):193-202. 8570901. *No comparison group*
48. Eames P, Cotterill G, Kneale TA, et al. Outcome of intensive rehabilitation after severe brain injury: a long-term follow-up study. *Brain Injury*. 1996 Sep;10(9):631-50. 8853867. *No comparison group*
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Appendix D

Appendix D. Table 1. Intermediate outcomes assessed in Included Studies

Intermediate Outcomes (more appropriate to rehabilitation settings or specific impairment domains)	Frequency
6-Item Interpersonal Support Evaluation List	1
Academic skills	1
Auditory Consonant Trigrams	1
Barthel Index (BI)	3
Bond Neurophysical Scale	1
Brief Symptom Inventory-18 (BSI-18)	1
BRISS-R, PDBS, and PCSS	1
Buschke Selective Reminding Test	1
California Verbal Learning Test II	3
Cambridge Prospective Memory Test (CAMPROMPT)	1
Center for Epidemiological Studies Depression Scale	1
Comprehensive Assessment of Prospective Memory (CAPM)	1
Control Oral Word Association	2
COWAT	1
Current Status-Relative Ratings (SPRS-Relative)	1
Depression Anxiety Stress Scales (DASS)	2
Facial Expression Matching Task	1
Facial Expression Naming Task	1
Functional Assessment Measure (FAM)	2
Functional Independence Measure (FIM)	5
Functional Status Examination	1
General Health Questionnaire	1
Glasgow Assessment Schedule	1
Glasgow Outcome Scale –Extended (GOS-E)	2
Goal Attainment Scaling (GAS)	1
Halstead-Reitan Neuropsychological Impairment Index	2
Higher order and conceptual skills	1

Intermediate Outcomes (more appropriate to rehabilitation settings or specific impairment domains)	Frequency
Katz Adjustment scale (KAS)	2
Katz Adjustment Scale-Relative's Form (KAS-R)	2
La Trobe Communication Questionnaire	1
Leeds Depression Scale	1
Logical memory	1
Mill Hill Vocabulary	1
Modified Health and Activity Limitations Survey (HALS)	1
Neurobehavioral Functioning Inventory (NFI)	1
Neurobehavioral Rating Scale	1
Orientation Remedial Module (ORM)	1
PASAT	1
Perceived self-efficacy	1
Profile of Functional Impairment in Communication (PFIC)	1
Psychomotor dexterity	1
Purdue Pegboard	1
Rey Complex Figure Test	4
Russell-Neurenger Average Impairment Rating (AIR)	1
Scales of Cognitive Ability for TBI	1
Social Activity Interview	1
Social Communication Skills Questionnaire Adapted (SCSQ-A)	1
Social Performance Survey Schedule	1
SPSS-Positive and SPSS Negative	1
Symbol Digits Modalities Test	1
The Awareness of Social Inference Test (TASIT), parts 1, 2, and 3	3
The Booklet Category Test	3
Therapeutic Alliance	1
Time in therapy	2

Intermediate Outcomes (more appropriate to rehabilitation settings or specific impairment domains)	Frequency
Trahan Continuous Visual Memory Test	1
Trail Making Tests	3
UCLA Loneliness Scale	1
Visual processing skills	1
WAIS-R	1
WAIS-verbal	1
Weekly social activity data	1
Weschler Abbreviated Scale of Intelligence	1
Weschler Memory Scale III	2
Weschler Memory Scale Revised	1
Wisconsin Card Sorting Test	2
Woodcock-Johnson III	1

Note: This table lists the outcomes assessed in eligible studies that we classified as intermediate outcomes. The 23 eligible studies assessed over 50 different intermediate outcomes scales.

Appendix D. Table 2. Secondary outcomes

Study, Design; Instrument	Treatment Arms	Outcome Before Treatment	Outcome After Completion of Treatment	Treatment vs. Control; Comments
Cicerone 2008,¹ RCT <u>Perceived Quality of Life (PQOL)</u> post treatment (16 weeks)	Intensive Cognitive Rehabilitation Program (ICRP) (n=34)	59.0 (21.7)	66.8 (17.5) P<0.05 versus before treatment	ES=0.26 [-0.22 to 0.74] No significant differences between groups but Intensive cognitive rehabilitation participants showed greater improvements on the PQOL
	Standard Neurorehabilitation Program (STD) (n=34)	61.2 (16.5)	62.2 (17.2)	
Vanderploeg 2008,² RCT <u>Disability Rating Scale (DRS)</u> 1 year post protocol treatment	Functional-experimental (n=150)	NR	8.2 (5.3)	ES=0.12 [-0.11 to 0.34] No significant differences between groups (P=0.29)
	Cognitive-didactic (n=152)	NR	7.6 (4.8)	
Vanderploeg 2008,² RCT <u>Quality of Life (satisfied with life- yes/no)</u> 1 year post protocol treatment	Functional-experimental (n=124)	NR	65% (81/124)	RR = 1.06 [0.88 to 1.28] No significant differences between groups (P=0.53)
	Cognitive-didactic (n=130)	NR	62% (80/130)	
Powell 2002,¹⁴ RCT <u>Brain injury community rehabilitation outcome- 39 (BICRO-39)</u> 27 weeks post treatment	Outreach (n=35 of 54 randomized)	Median (range) 15.3 (8 to 22.3)	% improving 80.0 (28/35) Median change (range) 2.5 (-1.7 to 6.2)	RR = 1.14 [0.88 to 1.49] Total BICRO-39 change score (summed across the six scales) was significantly greater in the outreach group than in the information group (mean ranks: outreach 43.2, information 33.4; U=517, p=0.05).
	Information (n=40 of 56 randomized)	Median (range) 12.9 (8.8 to 25.7)	% improving 70.0 (28/40) Median change (range) 0.9 (-4.1 to 6.8)	
Bell 2005¹³ RCT EuroQoL	Telephone	NR	Adjusted mean 0.78	Treatment effect=0.10 (0.02-0.19)
	Standard	NR	Adjusted mean 0.67	
Bell 2005¹³	Telephone	NR	Adjusted mean	Treatment effect=0.40 (-0.05-0.84)

Study, Design; <u>Instrument</u>	Treatment Arms	Outcome Before Treatment	Outcome After Completion of Treatment	Treatment vs. Control; Comments
RCT			6.58	
GOS-E	Standard	NR	Adjusted mean 6.19	
Bell 2005¹³	Telephone	NR	Adjusted mean 78.9	Treatment effect=8.8 (1.7-15.9)
RCT	Standard	NR	Adjusted mean 70.1	
PQoL				
Cicerone 2004¹¹	Intensive Cognitive Rehabilitation Program (ICRP) (n=34)	NR	27.1 (4.6)	Standard treatment group reported higher QCI scores (P<.01)
QCI	Standard Neurorehabilitation Program (STD) (n=34)	NR	29.7 (4.4)	
Thomas 2004¹⁵	Potential Unlimited Program	35.36 (8.80)	Stage 1 42.57 (11.08) Posttreatment 38.26 (10.56) 6-month followup 46.14 (12.22) 2-year followup 50.00 (13.95)	Only significant difference between groups at 6-month followup.
	No treatment	38.63 (21.97)	Stage 1 39.63 (19.66) Posttreatment 39.00 (18.88) 6-month followup 20.25 (14.73) 2-year followup 41.83 (10.36)	
Semlyen 1998¹⁶ quasi-experimental (CCT)	Multidisciplinary rehabilitation service (n=33)	Group differences in change 8 wk to 12 wk 4.00 (p<0.001)†	Group differences in change 6 mo to 12 mo 3.82 (p<0.01)†	The multidisciplinary rehabilitation service group showed significant gains throughout the rehabilitation period, the single discipline approach group did not.
Newcastle Independence Assessment Form (NIAF) 6-12 months post	Single discipline approach (n=18)	Group differences in change 8 wk to 12 wk 2.30 (p<0.05)†	Group differences in change 6 mo to 12 mo 1.05 (p NS)	

Study, Design; <u>Instrument</u>	Treatment Arms	Outcome Before Treatment	Outcome After Completion of Treatment	Treatment vs. Control; Comments
treatment (rehab period)				
Greenwood 1994⁴ GOS-E	Case-management (N=53 at entry)	NR	6 months posttreatment 5.3 (1.7) N=48 12 months posttreatment 5.5 (1.6) N=37 24 months posttreatment 5.6 (1.5) N=21	No group differences.
	Control (N=65 at entry)	NR	6 months posttreatment 5.8 (1.5) N=59 12 months posttreatment 6.2 (1.4) N=55 24 months posttreatment 6.3 (1.2) N=29	
Greenwood 1994⁴ GOS-E	Case-management (N=53 at entry)	NR	24 months posttreatment 2.0 (2.4) N=19	Case managed have significantly worse DRS scores. (p<0.05)
	Control (N=65 at entry)	NR	24 months posttreatment 0.6 (1.7) N=29	

* Based on Cohen's "Rules-of-Thumb" standardized mean difference effect size are as follows: small = 0.20; medium = 0.50; and large = 0.80. ** 25th and 75th quartiles. † For within group differences between means at each time point

ES = effect size; NS = not statistically significant; RR = Risk ratio [95% confidence interval]

Note: This table presents the results of studies that assessed a secondary outcome.

Appendix E

Appendix E. Table 1. Evidence table of multidisciplinary postacute rehabilitation for moderate to severe TBI studies

Study Description	Inclusion/Exclusion Criteria	Demographic/ Preinjury Characteristics	TBI Characteristics	Postinjury Characteristics
Bell, 2005¹³ Moderate to Severe TBI	Telephone Counseling Theory/Model: Modeled after validated telephone interventions in chronic care, smoking cessation, depression Program Type: Post-rehabilitation telephone contact Setting: Patient home Delivery: Scheduled phone calls with individualized mail supplements	Description: Scheduled phone calls made "research care manager to randomly allocated post-rehabilitation discharge patients. Calls were comprised of 3 basic elements: Follow-up of previously stated concerns, patient or family member stated current concerns, research care manager determined level of intervention in response to patient's concern. Coordination: NR Disciplines: NR Components: Giving information, mentoring, goal-setting, reassurance, modeling problem-solving, referral to community resources, triaging to regional or tertiary center if local resources unavailable Therapy hours/week: 30-45 minutes, weeks 2, 4 and months 2, 3, 5, 7, and 9 post-rehabilitation Duration: 9 months Total therapy hours: NR Manualized: Yes, described in detail in previous publication Staff Training: NR Fidelity Checks: NR	Bell, et al, 2005 [15895327] Moderate to Severe TBI	Telephone Counseling Theory/Model: Modeled after validated telephone interventions in chronic care, smoking cessation, depression Program Type: Post-rehabilitation telephone contact Setting: Patient home Delivery: Scheduled phone calls with individualized mail supplements

Appendix E. Table 1. Evidence table of multidisciplinary postacute rehabilitation for moderate to severe TBI studies

Study Description	Inclusion/Exclusion Criteria	Demographic/ Preinjury Characteristics	TBI Characteristics	Postinjury Characteristics
Cicerone, 2004¹¹	Inclusion criteria	Age (years±SD)	Severity (% moderate/severe)	Comorbidities
Study design Prospective Cohort	<ul style="list-style-type: none"> medically stable independent in basic self-care skills cognitive ability to participate in treatment medical documentation TBI 18 or older adequate language expression and comprehension 	<ul style="list-style-type: none"> ICRP 38±10.6 SRP 37±12.0 	<ul style="list-style-type: none"> ICRP 89% SRP 90% 	Psychiatric comorbidities not described, although subjects identified with current substance use or psychiatric disturbance that would preclude effective treatment for their cognitive deficits were not admitted. Psychiatric subjects were guided to the intensive cognitive group.
Sample size 57	Exclusion criteria	Gender (% male)	Severity definition NR	
Location Edison, NJ	<ul style="list-style-type: none"> current substance use or psychiatric disturbance precluding effective treatment no available family member or person to participate in program 	<ul style="list-style-type: none"> ICRP 63% SRP 79% 	Time since injury (months±SD)	
Setting Community-based, postacute outpatient brain injury rehabilitation program		Race/ethnicity NR	<ul style="list-style-type: none"> ICRP 33.9±4.8 SRP 4.8±9.5 	
Interventions		Education (years±SD)	TBI etiology NR	
<ul style="list-style-type: none"> Intensive cognitive rehabilitation group (ICRP) (n=27) (Control) Standard neurorehabilitation (SRP) (n=29) 		<ul style="list-style-type: none"> ICRP 13.2±1.7 SRP 13.0±2.2 	Area of brain injured NR	Compensation seeking NR
Primary outcomes CIQ		Employment status (% competitively employed)	Other injury characteristics NR	Acute rehabilitation history NR
		<ul style="list-style-type: none"> ICRP 96 SRP 97 		
		Income NR		
		Marital status NR		
		Military/Veteran NR		
		Insurance status NR		
		Prior TBI NR		
		Preexisting psychiatric conditions NR		Concomitant treatment NR

Appendix E. Table 1. Evidence table of multidisciplinary postacute rehabilitation for moderate to severe TBI studies

Study Description	Inclusion/Exclusion Criteria	Demographic/ Preinjury Characteristics	TBI Characteristics	Postinjury Characteristics
Cicerone, 2008¹	Inclusion Criteria:	Age (years, SD) ICRP: 39 (±11.) STD: 35 (±12.4)	Severity Mild: 13% Moderate: 24% Severe: 59%	Comorbidities: NR
Study design RCT	<ul style="list-style-type: none"> Medical documentation of TBI based on primary source within 24 hours of injury 	Gender (% male): 68%		Compensation seeking status: NR
Sample size 68	<ul style="list-style-type: none"> At least 3 months postinjury 		Severity Definition:	Acute rehabilitation history (% inpatient rehab)
Location Edison, NJ	<ul style="list-style-type: none"> 18-62 years of age Adequate language expression and comprehension (English) 	Race/ethnicity: 75% white, 10% black, 12% Hispanic, 3% Asian	Any combination of initial Glasgow Coma Scale score, duration of unconsciousness, duration of post-traumatic amnesia, and positive neuroimaging available from primary medical records.	ICRP: 77% STD: 85%
Setting Postacute brain injury rehabilitation center in suburban hospital	<ul style="list-style-type: none"> Judged to require at least 4 months comprehensive treatment Clinically appropriate for either arm of treatment Capable of attending treatment 3 days/week Capable of giving informed consent 	Education: (HS or <, some college, college grad)		Concomitant Treatment NR
Interventions	Exclusion Criteria:	Employment status: 79% employed, 4% unemployed, 2% homemaker, 13% student, 2% retired	Time since injury (mos mean, (std dev.)) ICRP=49.6 (±76.5) STD=37.0 (±58.2)	
<ul style="list-style-type: none"> Intensive cognitive rehabilitation (ICRP) Standard neurorehabilitation (STD) 	<ul style="list-style-type: none"> Active psychiatric illness, substance abuse, or pain that may prevent compliance with treatment 	Income: NR		
Primary Outcomes		Marital status(% married): 35%	TBI Etiology NR	
<ul style="list-style-type: none"> CIQ Vocational Integration Scale (community-based employment) 		Military/Veteran status: NR		
Secondary Outcomes		Insurance status: NR	Brain area injured NR	
<ul style="list-style-type: none"> Perceived Quality of Life scale (PQOL) 		Prior TBI: 4%	Other injury characteristics: NR	
		Preexisting psychiatric conditions: psychiatric illness 13% substance abuse 21%		

Appendix E. Table 1. Evidence table of multidisciplinary postacute rehabilitation for moderate to severe TBI studies

Study Description	Inclusion/Exclusion Criteria	Demographic/ Preinjury Characteristics	TBI Characteristics	Postinjury Characteristics
Greenwood, 1994⁴				
Study design prospective controlled unmatched nonrandomized study	Inclusion criteria <ul style="list-style-type: none"> closed head injury aged 16-60 been in coma for 6 hours or had a PTA > 48 hours care giver was resident in district informed consent Exclusion criteria <ul style="list-style-type: none"> received hospital treatment for drug or alcohol misuse aged 16-60 psychiatric disturbance, or a disorder of the central nervous system during the previous year no fixed abode or if follow up unlikely 	Age (years±SD) <ul style="list-style-type: none"> CM 31.6±14.4 control 30.7±14.0 Gender (% male) <ul style="list-style-type: none"> CM 69.6 control 75.7 Race/ethnicity NR Education NR Employment status (%) <ul style="list-style-type: none"> CM 100 control 96 Income NR Marital status NR Military/Veteran NR Insurance status NR Prior TBI NR Preexisting psychiatric conditions alcohol intake at injury (%) <ul style="list-style-type: none"> CM 36 control 37 	Severity definition “severely head injured patients” Severity GCS (mean±SD) <ul style="list-style-type: none"> CM 5.5±2.6 control 6.6±3.0 Duration of PTA (days±SD) <ul style="list-style-type: none"> CM 64.9±97.5 control 40.8±75.0 Time since injury NR TBI etiology (%) traffic accident/assault/fall/other <ul style="list-style-type: none"> CM <ul style="list-style-type: none"> traffic accident 60 assault 16 fall 18 other 5 control <ul style="list-style-type: none"> traffic accident 63 assault 14 fall 16 other 7 Area of brain injured NR MRI/imaging findings NR Other injury characteristics days unconscious (mean±SD) <ul style="list-style-type: none"> CM 11.3±13.5 control 4.6±7.5 	Comorbidities <ul style="list-style-type: none"> respiratory <ul style="list-style-type: none"> CM 47 control 21 conservative management <ul style="list-style-type: none"> CM 16 control 31 tracheostomy <ul style="list-style-type: none"> CM 32 control 16 Compensation seeking (%) <ul style="list-style-type: none"> 6 months <ul style="list-style-type: none"> CM 2 control 2 12 months <ul style="list-style-type: none"> CM 0 control 6 24 months <ul style="list-style-type: none"> CM 17 control 4 Acute rehabilitation history NR Concomitant treatment NR
Sample size 126 (outcomes for 118)				
Location four district general hospitals and two university teaching hospitals with neurosurgical units				
Setting London and environs				
Interventions <ul style="list-style-type: none"> case managed (CM) (n=56) control (n=70) 				
Secondary outcomes <ul style="list-style-type: none"> DRS GOS 				

Appendix E. Table 1. Evidence table of multidisciplinary postacute rehabilitation for moderate to severe TBI studies

Study Description	Inclusion/Exclusion Criteria	Demographic/ Preinjury Characteristics	TBI Characteristics	Postinjury Characteristics
Hashimoto 2006¹⁰	Inclusion criteria	Age (years±SD)	Severity definition	Comorbidities
Study design prospective, nonrandomized controlled trial	<ul style="list-style-type: none"> • near independence in Activities of Daily Living (ADL) irrespective of ability to walk or wheelchair use • the goal of returning to work or school • having no place they were required to visit frequently except for outpatient clinic 	<ul style="list-style-type: none"> • intervention 26.6±9.7 • control 28.7±10.9 	GCS ≤ 8	NR
Sample size 37		Gender (% male)	Severity (%)	Compensation seeking
Location Kanagawa Prefecture, Japan	Exclusion criteria NR	<ul style="list-style-type: none"> • intervention 72 • control NR 	<ul style="list-style-type: none"> • intervention 76.0 • control 83.3 	NR
Setting Kanagawa Rehabilitation Hospital		Race/ethnicity NR	Duration of PTA NR	Acute rehabilitation history NR
Interventions		Education NR	Time since injury (days±SD)	Concomitant treatment
<ul style="list-style-type: none"> • comprehensive day treatment program (n=25) • control (outpatients with TBI) (n=12) 		Employment status (% competitively employed)	<ul style="list-style-type: none"> • intervention 527.3±512.6 • control 487.6±125.9 	NR
Primary outcomes		<ul style="list-style-type: none"> • intervention 60 • control NR 	TBI etiology (%)	
<ul style="list-style-type: none"> • return to work • FIM/FAM • CIQ 		Income NR	<ul style="list-style-type: none"> • intervention <ul style="list-style-type: none"> ○ auto accident 20 ○ pedestrian/auto 20 ○ bike/auto 36 ○ cerebral aneurysm 8 ○ glioma 4 ○ fall 8 ○ work accident 4 • control NR 	
		Marital status NR	Area of brain injured	
		Military/Veteran NR	<ul style="list-style-type: none"> • intervention <ul style="list-style-type: none"> ○ diffuse brain injury 64 ○ diffuse brain injury + right acute subdural hematoma 20 ○ right acute subdural hematoma 4 ○ Sub arachnoid hemorrhage 8 ○ diffuse brain injury + contusion 4 • control NR 	
		Insurance status NR		
		Prior TBI NR		
		Pre-existing psychiatric conditions NR		
			MRI/imaging findings NR	

Appendix E. Table 1. Evidence table of multidisciplinary postacute rehabilitation for moderate to severe TBI studies

Study Description	Inclusion/Exclusion Criteria	Demographic/ Preinjury Characteristics	TBI Characteristics	Postinjury Characteristics
Ponsford, 2006⁵	Inclusion criteria Moderate to severe TBI patients	Age (years±SD) <ul style="list-style-type: none"> Community based 35.43±16.65 Control 33.78±15.41 	Severity (mean GCS±SD) <ul style="list-style-type: none"> Community based 8.22±4.37 Control 7.76±4.13 	Comorbidities NR
Study design Controlled, individually matched cohort trial	Exclusion criteria NR	Gender (% male) <ul style="list-style-type: none"> Community based 73 Control 73 	Severity definition GCS	Compensation seeking NR
Sample size 77		Race/ethnicity NR	Time since injury (years) NR	Acute rehabilitation history NR
Location Melbourne, Australia		Education (years±SD) <ul style="list-style-type: none"> Community based 11.56±2.42 Control 11.15±2.54 	TBI Etiology NR	Concomitant treatment NR
Setting Rehabilitation center		Employment status (% competitively employed) <ul style="list-style-type: none"> Community based 66 Control 70 	Area of brain injured NR	
Interventions <ul style="list-style-type: none"> Community based rehabilitation (n=77) Control (n=77) 		Income NR	Other injury characteristics NR	
Primary outcomes Return to work		Marital status (% single) <ul style="list-style-type: none"> Community based 63 Control 61 		
		Military/Veteran NR		
		Insurance status NR		
		Prior TBI NR		
		Preexisting psychiatric conditions NR		

Appendix E. Table 1. Evidence table of multidisciplinary postacute rehabilitation for moderate to severe TBI studies

Study Description	Inclusion/Exclusion Criteria	Demographic/ Preinjury Characteristics	TBI Characteristics	Postinjury Characteristics
Powell, 2002¹⁴	Time since injury (yrs mean, (std dev.)): Outreach=4.0±4.9, Information=2.7±3.6	Age (years, SD) Outreach=34±11, Information=35±10	Severity Mild: 1% Moderate: 0% Severe: 99%	Comorbidities NR
Study design: RCT				Compensation seeking status NR
Sample size 94	Inclusion Criteria:	Gender (% male): 76%		
Location London, England	<ul style="list-style-type: none"> • Age 16-65 • Severe TBI between 3 months and 20 years previously • No other neurological conditions • Reside within 1 hour travel time of hospital • Long-term treatment goals amenable with intervention 	Race/ethnicity NR	Severity Definition: Severe: PTA >1day Mild: PTA ≤ 1 hour	Social support: NR
Setting Community-based		Education NR	TBI Etiology NR	Acute rehabilitation history: community or post-rehab discharge
Study design: RCT		Employment status NR	Brain area injured NR	Concomitant Treatment NR
Interventions:	Exclusion Criteria NR	Income NR	Other injury characteristics: NR	
<ul style="list-style-type: none"> • Outreach • Information 		Marital status NR		
Primary Outcomes		Military/Veteran status NR		
<ul style="list-style-type: none"> • none 		Insurance status NR		
Secondary Outcomes		Prior TBI NR		
<ul style="list-style-type: none"> • BICRO-39 • 		Psychiatric conditions NR		
Intermediate Outcomes				
<ul style="list-style-type: none"> • BICRO-39 • FIM + FAM 				

Appendix E. Table 1. Evidence table of multidisciplinary postacute rehabilitation for moderate to severe TBI studies

Study Description	Inclusion/Exclusion Criteria	Demographic/ Preinjury Characteristics	TBI Characteristics	Postinjury Characteristics
Prigatano, 1984⁹	Inclusion criteria NR	Age (years±SD) <ul style="list-style-type: none">• Neuropsychologic 26.1±8.3• Control NR	Severity (% moderate/severe) NR	Comorbidities NR
Study design retrospective, controlled cohort study	Exclusion criteria NR	Gender (% male) <ul style="list-style-type: none">• Neuropsychologic 83.3• Control NR	Severity Definition Russell-Neurenger Average Impairment Rating	Compensation seeking NR
Sample size 18		Race/ethnicity NR	Time since injury (months) <ul style="list-style-type: none">• Neuropsychologic 21.6• Control NR	Acute rehabilitation history NR
Location Oklahoma City, Oklahoma		Education (%) <ul style="list-style-type: none">• Neuropsychologic<ul style="list-style-type: none">◦ ≤ 12 years 61.1◦ > 12 years 38.9• Control NR	TBI etiology “Severe closed head injury”	Concomitant treatment NR
Setting Neuropsychological rehabilitation program		Employment status (% competitively employed) <ul style="list-style-type: none">• Neuropsychologic 72.2• Control NR	Area of brain injured (%) <ul style="list-style-type: none">• Neuropsychologic<ul style="list-style-type: none">◦ Severe cerebral contusion 61.1◦ Brain stem contusion 5.6◦ Severe cerebral contusion + brain stem contusion 33.3• Control NR	
Interventions <ul style="list-style-type: none">• Psychotherapeutic (n=18)• Control (n=18)		Income NR	Other injury characteristics (%) <ul style="list-style-type: none">• Neuropsychologic<ul style="list-style-type: none">◦ Post traumatic seizure disorder 16.7◦ Residual paresis 66.7◦ Residual signs of aphasia and/or dysarthria 33.3◦ “Virtually all . . . had cerebral contusions and/or brain stem contusion”• Control NR	
Primary outcomes Return to work		Marital status NR		
		Military/Veteran (%) <ul style="list-style-type: none">• Neuropsychologic 5.6• Control NR		
		Insurance status NR		
		Prior TBI NR		
		Preexisting psychiatric conditions NR		

Appendix E. Table 1. Evidence table of multidisciplinary postacute rehabilitation for moderate to severe TBI studies

Study Description	Inclusion/Exclusion Criteria	Demographic/ Preinjury Characteristics	TBI Characteristics	Postinjury Characteristics
Prigatano, 1994⁷	Inclusion criteria	Age (years±SD)	Severity (mean±SD)	Comorbidities
Study design Matched control, prospective cohort	<ul style="list-style-type: none"> Primary diagnosis of craniocerebral trauma or TBI By end of study, ≥ 15 months elapsed since injury Admitted to study 2-55 months from injury All subjects considered potentially able to return to work/school 	<ul style="list-style-type: none"> Neuropsychological rehab 29.6±12.7 Historic controls (28.7±12.2) 	<ul style="list-style-type: none"> Neuropsychological rehab 8.08±2.7 Historic controls (n=38) 8.03±2.8 	NR
Sample size 79 (outcomes for 76)		Gender (% male)	Severity definition GCS	Compensation seeking NR
Location Phoenix, Arizona	Exclusion criteria NR	Race/ethnicity NR	Time since injury (months±SD)	Acute rehabilitation history NR
Setting Work Re-entry Program of the Adult Day Hospital for Neurological Rehabilitation at Saint Joseph's Hospital		Education (years±SD)	<ul style="list-style-type: none"> Neuropsychological rehab 43.3±16.1 Historic controls 33.5±8.7 	Concomitant treatment NR
Interventions		Employment status (% competitively employed)	TBI etiology NR	
<ul style="list-style-type: none"> Neuropsychological rehab (n=41, outcomes for 38) Historic controls (n=38) 		<ul style="list-style-type: none"> Neuropsychological rehab 78.0 Historic controls NR 	Area of brain injured NR	
Primary outcomes Return to work		Income NR	Other injury characteristics (%)	
		Marital status NR	<ul style="list-style-type: none"> Neuropsychological rehab <ul style="list-style-type: none"> CT/MRI findings of contusion and/or hematoma 87.7 Skull fracture/no hematoma 4.9 Loss of consciousness 7.3 Historic controls NR 	
		Military/Veteran NR		
		Insurance status NR		
		Prior TBI NR		
		Preexisting psychiatric conditions NR		

Appendix E. Table 1. Evidence table of multidisciplinary postacute rehabilitation for moderate to severe TBI studies

Study Description	Inclusion/Exclusion Criteria	Demographic/ Preinjury Characteristics	TBI Characteristics	Postinjury Characteristics
Rattok, 1992⁸	Inclusion Criteria	Age (median years)	Severity definition	Prior psychiatric conditions (%)
Study design 3 group comparison	<ul style="list-style-type: none"> • Diagnosis of TBI, ≥1hr coma • Diagnosis of cerebral anoxia, ≥12hr coma • ≥1 year post-injury • Neurological stability • Unsuccessful vocational or educational rehabilitation prior to entry into program • Residence in New York metropolitan area for duration of study 	<ul style="list-style-type: none"> • Treatment 1: 26.8 • Treatment 2: 27.1 • Treatment 3: 28.5 	Severe=Coma of ≥1hr or cerebral anoxia of ≥12hrs	<ul style="list-style-type: none"> • NR
Sample size 59	<ul style="list-style-type: none"> • Age, 18-55 • Command of English • Partial independence in basic activities of self-care, ambulation, and continence • Minimum IQ of 80 on WAIS • Minimum motivation for rehabilitation • Basic level of social appropriateness and manageability in therapeutic or training environment 	Gender (% male)	Severity (Days in coma)	Comorbidities (%)
Location New York, NY Metropolitan Area		<ul style="list-style-type: none"> • Treatment 1: 65% • Treatment 2: 89% • Treatment 3: 61% 	<ul style="list-style-type: none"> • Treatment 1: 34.3 • Treatment 2: 38.9 • Treatment 3: 36.9 	<ul style="list-style-type: none"> • NR
Setting Outpatient rehabilitation center		Race/ethnicity (%) NR	Time since injury (median months)	Compensation seeking NR
Interventions		Education (median years)	<ul style="list-style-type: none"> • Treatment 1: 32 • Treatment 2: 33.8 • Treatment 3: 40.2 	Acute rehabilitation history “Unsuccessful”
<ul style="list-style-type: none"> • Treatment 1 (Balanced) • Treatment 2 (Interpersonal) • Treatment 3 (Individualized) 	Exclusion criteria	Employment status (% competitively employed) NR	TBI etiology 95% acceleration/deceleration concussion; 5% cerebral anoxia	
Primary outcomes	<ul style="list-style-type: none"> • History or present psychiatric complications • History of drug or alcohol abuse • History of sociopathy • Inability to communicate 	Income NR	MRI/imaging findings NR	
<ul style="list-style-type: none"> • Cognitive performance measures • Behavioral Competence Index (BCI) • Vocational 		Marital status (%) NR	Other injury characteristics (%)	<ul style="list-style-type: none"> • NR
		Military/Veteran NR		
		Insurance status NR		
		Prior TBI (%) NR		

Appendix E. Table 1. Evidence table of multidisciplinary postacute rehabilitation for moderate to severe TBI studies

Study Description	Inclusion/Exclusion Criteria	Demographic/ Preinjury Characteristics	TBI Characteristics	Postinjury Characteristics
Salazar, 2000³ Study design: RCT Sample size 120 Location: Washington, D.C. Setting US Military medical referral center Interventions: <ul style="list-style-type: none"> Intensive, interdisciplinary, in-hospital cognitive rehabilitation program (Hospital) (n=xx) Limited home rehabilitation program with telephone support from psychiatric nurse (Home) (n=xx) Primary Outcomes <ul style="list-style-type: none"> Return to work Fitness for military duty Secondary Outcomes <ul style="list-style-type: none"> none 	Inclusion Criteria: <ul style="list-style-type: none"> Moderate-to-severe closed head injury Head injury within 3 months of randomization Rancho Los Amigos cognitive level of 7 Active duty military member; not pending separation Accompanied home setting with at least 1 responsible adult available Ability to independently ambulate No prior severe TBI or other severe disability that would preclude return to active duty after study treatment Exclusion Criteria: <ul style="list-style-type: none"> Mild TBI 	Age: Hospital=25, 6.63; Home=26,6.22 Gender(% male): Hospital: 93% Home: 96% Race/ethnicity(% white) Hospital: 69% Home: 70% Education (% some college): Hospital: 41% Home=44% Employment status: NR Income: NR Marital status (% married) Hospital:30% Home=34% yes Military/Veteran status(% active military): 100% Insurance status (% military coverage): 100% Prior TBI Hospital: 11% Home: 18% Psychiatric conditions(% positive diagnosis) Hospital=19% Home=25%	Severity Severity Definition Glasgow Coma Scores≤13; or posttraumatic amnesia≥24 hours; or focal cerebral contusion or hemorrhage on computed tomography or MRI Time since injury (mean days, SD) Hospital: 38 (23.6) Home: 39 (33.2) Etiology MVC Hospital:49% Home: 72% Assault: Hospital: 27% Home: 9% Unknown: Hospital: 24% Home: 19% Area of brain injured: cerebrum; computed tomography or MRI Other injury characteristics Closed: 100%	Comorbidities: Headaches, violent behavior, aggressive behavior, seizures, major depression Compensation seeking status: NR Social support: Accompanied home setting with at least 1 responsible adult available Acute rehabilitation history: NR Concomitant Treatment NR

Appendix E. Table 1. Evidence table of multidisciplinary postacute rehabilitation for moderate to severe TBI studies

Study Description	Inclusion/Exclusion Criteria	Demographic/ Preinjury Characteristics	TBI Characteristics	Postinjury Characteristics
Sarajuuri, 2005⁶	Inclusion Criteria	Age (at injury; years, SD)	Severity (admission GCS; mean, SD, range)	Comorbidities: NR
Study design Prospective Cohort	<ul style="list-style-type: none"> Independence in daily life and only slight physical disabilities 16 to 55 years of age completed compulsory education adequate potential to achieve productivity 	T: 30.5 (\pm 10.6) C: 29.5 (\pm 11.0)	T: 7.9 (2.7) (4-14) C: 8.2 (2.5) (3-13)	Compensation seeking NR
Sample size 39		Gender (% male)	Severity Definition:	Acute rehabilitation history
Location Helsinki, Finland	Exclusion Criteria	T: 84% C: 85%	NR	OT T: 32% C: NR
Setting Nationwide Rehabilitation Center & Neurosurgery Department within academic medical center hospital	<ul style="list-style-type: none"> significant psychiatric history alcohol or drug abuse previous brain injury another malignant disease 	Race/ethnicity NR	Time since injury (month,SD)	PT T: 47% C: NR
Interventions	Population (n)	Education (years, SD)	T: 84% C: 85%	SLP T: 26% C: NR
<ul style="list-style-type: none"> Comprehensive (T) (n=19) Conventional (C) (n=20) 	T: 19 C: 20	Employment status (preinjury; % employed or studying preinjury)	TBI Etiology (% by mechanism)	NP T: 37% C: NR
Primary Outcome Status of productivity		T: 84% C: 85%	MVC/bike/pedestrian T: 63% C: 55%	Concomitant Treatment NR
		Income NR	Assault T: 5% C: 5%	
		Marital status NR	Other(fall, hit by) T: 26% C: 40%	
		Military/Veteran NR	Unknown T: 5% C: 0%	
		Insurance status NR	Area of brain injured:	
		Prior TBI NR, but prior TBI is excluded	NR	
		Preexisting psychiatric conditions NR, but significant psychiatric history excluded	Other Injury characteristics	
			Contusion/hematoma T: 79% C: 80%	
			Diffuse axonal injury T: 42% C: 25%	
			Severe intracranial pressure T: 37% C: 25%	
			Craniotomy T: 21% C: 25%	

Appendix E. Table 1. Evidence table of multidisciplinary postacute rehabilitation for moderate to severe TBI studies

Study Description	Inclusion/Exclusion Criteria	Demographic/ Preinjury Characteristics	TBI Characteristics	Postinjury Characteristics
Semlyen, 1998¹⁶	Inclusion Criteria:	Age (at injury; years, SD)	Severity	Comorbidities: NR
Study design	<ul style="list-style-type: none"> Initial Glasgow Coma Scale scores ≤ 8 for at least 6 hours 	Treatment: 36(13) Control: 30(12)	Severe: 100%	
Prospective Cohort	<ul style="list-style-type: none"> Between 16-65 years Identifiable primary consentor Resides in Northern Regional Health Authority Surgically stable and able to be discharged from neurosurgical unit within 4 weeks of injury 	Gender (% male)	Severity Definition	Compensation seeking status: NR
Sample size 51		Treatment: 85% Control: 84%	Severe: GCS Score ≤ 8 for at least 6 hours	Acute rehabilitation history: NR
Location: Newcastle upon Tyne, UK		Race/ethnicity: NR	Time since injury (mean days, SD)	Concomitant Treatment NR
Setting Regional rehabilitation centre	Exclusion Criteria:	Education: NR	Treatment: 49.37 (29.62) Control: 17.94 (13.6)	
	<ul style="list-style-type: none"> Previous drug or alcohol misuse Premorbid neurologic history 	Employment status: NR	TBI Etiology	
Interventions:		Income	<u>MVC</u>	
<ul style="list-style-type: none"> Coordinated, multidisciplinary rehabilitation Single-discipline rehabilitation 		"majority in both groups in lower-middle SES"	Treatment: 69.8% Control: 44.6%	
Primary Outcomes		Marital status: NR	<u>Falls</u>	
<ul style="list-style-type: none"> None 		Military/Veteran status: NR	Treatment: 18.2% Control: 33.3%	
Secondary Outcomes		Insurance status: NR	<u>Assault</u>	
<ul style="list-style-type: none"> Newcastle Independence Assessment Form-Research (NIAF-R) 		Prior TBI: NR	Treatment: 9.1% Control: 22.2%	
Intermediate Outcomes		Psychiatric conditions: NR	<u>Self-harm</u>	
<ul style="list-style-type: none"> Barthel Index FIM 			Treatment: Control: 3%	
			Brain area injured: NR	
			Other injury characteristics NR	

Appendix E. Table 1. Evidence table of multidisciplinary postacute rehabilitation for moderate to severe TBI studies

Study Description	Inclusion/Exclusion Criteria	Demographic/ Preinjury Characteristics	TBI Characteristics	Postinjury Characteristics
Thomas, 2004¹⁵	Inclusion Criteria <ul style="list-style-type: none"> Self-selected volunteers ABI Past or present client of NSW Brain Injury Rehabilitation Programme Exclusion criteria <ul style="list-style-type: none"> NR 	Age (mean years±SD) <ul style="list-style-type: none"> PUP 31.54±10.37 Controls 38.38±12.14 Gender (% male) <ul style="list-style-type: none"> PUP NR Control NR Race/ethnicity (%) <ul style="list-style-type: none"> PUP NR Control NR Education (mean years±SD) <ul style="list-style-type: none"> Intensive therapy 13.2±1.9 Standard therapy 12.5±1.2 Employment status (% competitively employed) <ul style="list-style-type: none"> PUP “Most not working/studying” Control “Most not working/studying” Income <p>NR</p> Marital status (%) <ul style="list-style-type: none"> PUP NR Control NR Military/Veteran <p>NR</p> Insurance status <p>NR</p> Prior TBI (%) <ul style="list-style-type: none"> PUP NR Control NR 	Severity definition <p>Mild=PTA 5-60 minutes Severe=PTA 1-7 days Very Severe=PTA 7-28 days Extremely Severe=PTA>28 days</p> Severity (%) <ul style="list-style-type: none"> PUP <ul style="list-style-type: none"> Mild 2 Severe 1 Very Severe 2 Extremely Severe 8 Control <ul style="list-style-type: none"> Mild 2 Severe 3 Very Severe 0 Extremely Severe 3 Time since injury (mean years±SD) <ul style="list-style-type: none"> PUP <ul style="list-style-type: none"> 5.99±4.54 Control <ul style="list-style-type: none"> 4.97±2.28 TBI etiology <p>NR</p> MRI/imaging findings <p>NR</p> Other injury characteristics (%) <ul style="list-style-type: none"> NR 	Prior psychiatric conditions (%) <ul style="list-style-type: none"> NR Comorbidities (%) <p>prior substance abuse</p> <ul style="list-style-type: none"> NR Compensation seeking <p>NR</p> Acute rehabilitation history (%) <p>All participants in PUP and control were current or past clients of New South Wales Brain Injury Rehabilitation Programme</p>

Appendix E. Table 1. Evidence table of multidisciplinary postacute rehabilitation for moderate to severe TBI studies

Study Description	Inclusion/Exclusion Criteria	Demographic/ Preinjury Characteristics	TBI Characteristics	Postinjury Characteristics
Vanderploeg, 2008² Study design RCT, Multicenter Sample size 366 Location Minneapolis, Palo Alto, Richmond, Tampa Setting VA acute inpatient TBI rehab programs Interventions <ul style="list-style-type: none"> Cognitive-didactic (CD) rehab therapy (n=184) Functional-experiential (FE) (n=182) Primary Outcomes <ul style="list-style-type: none"> Return to work Secondary Outcomes <ul style="list-style-type: none"> Disability Rating Scale score Functional independence in living 	<p>Inclusion Criteria: (1) moderate-to-severe nonpenetrating TBI within the preceding 6 months, manifested by a postresuscitation Glasgow Coma Scale score of 12 or less, or coma of 12 hours or more, or PTA of 24 hours or more, and/or focal cerebral contusion or hemorrhage on CT or MRI; (2) RLAS cognitive level of 5 to 7 at time of randomization; (3) age 18 years or older; (4) active duty military member or veteran; and (6) anticipated length of needed acute interdisciplinary TBI rehabilitation of 30 days or more</p> <p>Exclusion Criteria: (1) history of prior inpatient acute rehabilitation for the current TBI and (2) history of a prior moderate to severe TBI or other preinjury severe neurologic or psychiatric condition, such as psychosis, stroke, multiple sclerosis, or spinal cord injury</p>	<p>Age (at injury; years, SD) CD 33.2 (±13.5) FE 31.7 (±12.9)</p> <p>Gender (% male) CD: 92% FE:95%</p> <p>Race/ethnicity Hispanic CD: 10% FE:11% White CD: 68% FE:69% Black CD: 20% FE:18% Other CD: 12% FE:12%</p> <p>Education (% post high school) CD: 34% FE:37%</p> <p>Employment status: (% working or in school) CD: 86% FE:89%</p> <p>Income: NR</p> <p>Marital status (% married) CD: 25.6% FE: 25.1%</p> <p>Military/Veteran status (% what?) CD: 58.4% FE:67.8%</p> <p>Insurance status: NR</p> <p>Prior TBI (% “prior head injury”) CD: 7.2% FE: 7.2%</p> <p>Pre-existing psychiatric conditions: NR</p>	<p>Severity NR, but moderate/severe inclusion criteria</p> <p>Severity Definition: NR</p> <p>Time since injury:</p> <ul style="list-style-type: none"> CD 48.9±28.5 (n = 180) days FE 51.1±29.8 (n = 180) days <p>TBI Etiology: MVC CD: 68% FE:66% Assault CD: 10% FE:8%</p> <p>Area of brain injured: NR</p> <p>Injury characteristics:</p> <ul style="list-style-type: none"> CD <ul style="list-style-type: none"> Motor vehicular 122/180 (67.8%) Fall 21/180 (11.7%) Blunt object 15/180 (8.3%) Sports/training accident 5/180 (2.8%) Indeterminant 17/180 (9.4%) FE <ul style="list-style-type: none"> Motor vehicular 119/180 (66.1%) Fall 29/180 (16.1%) Blunt object 9/180 (5.0%) Sports/training accident 6/180 (3.3%) Indeterminant 17/180 (9.4%) 	<p>Comorbidities: NR</p> <p>Compensation seeking status: NR</p> <p>Acute rehabilitation history: NR</p> <p>Concomitant Treatment NR</p>

Appendix E. Table 1. Evidence table of multidisciplinary postacute rehabilitation for moderate to severe TBI studies

Study Description	Inclusion/Exclusion Criteria	Demographic/ Preinjury Characteristics	TBI Characteristics	Postinjury Characteristics
Willer, 1999¹²	Inclusion criteria Individuals with brain injury who had not undergone treatment in this community-based program	Age (years±SD) <ul style="list-style-type: none">• RBPR 33.42±11.31• Control 34.76±10.72	Severity (% moderate/severe) All subjects were considered severe TBI	Comorbidities NR
Study design Case controlled study using a matched design in a before-and-after trial	Exclusion criteria NR	Gender (% male) <ul style="list-style-type: none">• RBPR 87• Control 87	Severity Definition (HALS disability score±SD) <ul style="list-style-type: none">• RBPR 20.39±6.02• Control 20.30±6.09	Compensation seeking NR
Sample size 46		Race/ethnicity NR		Acute rehabilitation history NR
Location Ontario, Canada		Education (%) <ul style="list-style-type: none">• RBPR<ul style="list-style-type: none">◦ < HS 26.0◦ Completed HS 43.5◦ > HS 30.4• Control<ul style="list-style-type: none">◦ < HS 26.0◦ Completed HS 34.8◦ > HS 39.1	Time since injury (years±SD) <ul style="list-style-type: none">• RBPR 3.05±2.98• Control 4.66±4.66	Concomitant treatment NR
Setting Postacute residential rehabilitation program or home-based subjects			TBI etiology (%) <ul style="list-style-type: none">• RBPR<ul style="list-style-type: none">◦ Vehicular related 95.7◦ Assault 4.3• Control<ul style="list-style-type: none">◦ Vehicular related 95.7◦ Assault 4.3	
Interventions <ul style="list-style-type: none">• Residential-based postacute rehabilitation (RBPR) (n=23)• Control (n=23)		Employment status NR		
		Income NR		
		Marital status NR		
Primary outcomes CIQ		Military/Veteran NR	Area of brain injured NR	
		Insurance status NR		
		Prior TBI NR	Other injury characteristics Closed brain injury	
		Preexisting psychiatric conditions <ul style="list-style-type: none">• RBPR: 30.4% were recruited from psychiatric hospitals• Control NR		

Appendix E. Table 2. Intervention Characteristics

Study Target Population	Intervention Arm	Intervention Description and Implementation
Bell, 2005¹³ Moderate to Severe TBI	Telephone Counseling	<p>Description: Scheduled phone calls made “research care manager to randomly allocated post-rehabilitation discharge patients. Calls were comprised of 3 basic elements: Follow-up of previously stated concerns, patient or family member stated current concerns, research care manager determined level of intervention in response to patient’s concern.</p> <p>Coordination: NR</p> <p>Disciplines: NR</p> <p>Components: Giving information, mentoring, goal-setting, reassurance, modeling problem-solving, referral to community resources, triaging to regional or tertiary center if local resources unavailable</p> <p>Therapy hours/week: 30-45 minutes, weeks 2, 4 and months 2, 3, 5, 7, and 9 post-rehabilitation</p> <p>Duration: 9 months</p> <p>Total therapy hours: NR</p> <p>Manualized: Yes, described in detail in previous publication Staff Training: NR Fidelity Checks: NR</p>
	Standard Follow-up	<p>Description: Patient given recommendations from acute care team then not contacted until 1 year follow-up</p> <p>Coordination: NR</p> <p>Disciplines: primarily NR</p> <p>Components: NR</p> <p>Therapy hours/week: NR</p> <p>Duration: 1 year</p> <p>Total therapy hours: NR</p> <p>Manualized: NR Staff Training: NR Fidelity Checks: NR</p>

Appendix E. Table 2. Intervention Characteristics

Study Target Population	Intervention Arm	Intervention Description and Implementation
Cicerone, 2004 ¹¹ Chronic Moderate to Severe TBI	Intensive Cognitive Rehabilitation Program (ICRP) Theory/Model: Holistic neuropsychological rehabilitation (Ben-Yishay and Gold 1990) Program Type: Community-based day treatment program Setting: Suburban postacute brain injury rehabilitation center (US) Delivery: Peer groups progress through program together.	Description: 'Individual and group cognitive remediation with an emphasis on increasing awareness and developing compensations for cognitive deficits, small-group treatment for interpersonal and pragmatic communication skills, individual and/or group psychotherapy, family support, and therapeutic work trials and placement to facilitate educational or vocational readiness.' Coordination: NR Disciplines: NP, VT,; PT, OT if necessary Components: Cognitive group - 6 hrs/wk; individual cognitive remediation - 3 hrs/wk; communication and interpersonal skills group - 3 hrs/wk; applied skills group - 1 hr/wk; additional tailored therapies - variable/wk; therapeutic work trials – 1 day/wk; family involvement. Therapy hours/week: 15 hrs/wk Duration: 16 weeks Total therapy hours: 240 hours. Manualized: NR Staff Training: NR Fidelity Checks: NR
	Standard Rehabilitation Program (SRP) Theory/Model: 'conventional program' Program Type: Community-based day treatment program Setting: Postacute brain injury rehabilitation center (Suburban US) Delivery: Individuals progress through tailored treatments	Description: Treatment content and duration tailored to individual. Coordination: monitored by staff NP throughout course of treatment Disciplines: primarily NP, PT, OT, SLP; could also include RT, VT, E psychologic counseling Components: Tailored, typical patterns NR Therapy hours/week: 15 hrs/ wk initially, adjusted individually to range of 12 to 24 hr/ wk. Duration: 3.9 mo (mean) Total therapy hours: variable Manualized: NR Staff Training: NR Fidelity Checks: NR

Appendix E. Table 2. Intervention Characteristics

Study Target Population	Intervention Arm	Intervention Description and Implementation
Cicerone, 2008 ¹ Chronic Moderate to Severe TBI	Intensive Cognitive Rehabilitation Program (ICRP) Theory/Model: Berquist 1994; Holistic neuropsychological rehabilitation (Ben-Yishay and Gold 1990) Program Type: Day treatment program Setting: Suburban postacute brain injury rehabilitation center (US) Delivery: Peer groups progress through program together.	Description: Integrated treatments for cognitive deficits, interpersonal and behavioral difficulties, functional skills within therapeutic environment. Meta-cognition, emotional regulation, compensatory approaches emphasized. Weeks grouped by themes. Coordination: Disciplines: NP, primary therapist Components: Cognitive group - 6 hrs/wk; communication and interpersonal skills group - 3 hrs/wk; life skills group - 2 hr/wk; individual therapy - 3 hrs/wk, individual NP treatment 1 hr/wk. Therapy hours/week: 15 hr/wk Duration: 16 weeks Total therapy hours: 240 Manualized: NR Staff Training: NR Fidelity Checks: Yes
	Standard Neurorehabilitation Program (STD) Theory/Model: Comprehensive , interdisciplinary day treatment program (Malec 1996 Berquist 1994 Program Setting/Type: Day treatment program Setting: Postacute brain injury rehabilitation center (Suburban US) Delivery: Individuals progress through tailored treatments	Description: Individual, discipline-specific therapies targeting specific deficit areas designed to be responsive to stage and rate of recovery after TBI. Restorative strategies. Coordination: Followed by NP. Disciplines: NP, Psych, PT, OT, SLP, RT, VT, EC Components: Amounts and combinations of therapies varied. Most participants: individual NP treatment – 1 hr/wk; Participants could receive psychological counseling – 1 hr/wk, RT, VT, or educational counseling – 1 hr/wk; group therapy limited to 3 hrs/wk Therapy hours/week: 15 Duration: 16 weeks Total therapy hours: 240. Manualized: NR Staff Training: NR Fidelity Checks: Yes

Appendix E. Table 2. Intervention Characteristics

Study Target Population	Intervention Arm	Intervention Description and Implementation		
Greenwood, 1994⁴ Severe TBI	Case Management	Description: Early (within 7 days of injury) case management program which served as facilitator rather than therapeutic role, recruiting services for patient from a variety of agencies. Coordination: Case manager Disciplines: Physiotherapy, occupational therapy, speech therapy, psychology, social work Components: Determining patient needs and recruiting services based on these needs Therapy hours/week: NR Duration: NR; outcomes reported at 6, 12, and 24 months Total therapy hours: NR Manualized: Yes, described in detail in previous publication Staff Training: NR Fidelity Checks: NR		
	Theory/Model: Case management model established by authors in previous papers; “assertive” or “clinical” case management elements developed by Holloway for mentally ill Program Type: Pro-active case management Setting: 4 general hospitals and 2 university teaching hospitals Delivery: Home-based outreach			
	Control	Description: Patient given standard rehabilitation without case management Coordination: NR Disciplines: Physiotherapy, occupational therapy, speech therapy, psychology, social work Components: NR Therapy hours/week: NR Duration: NR, outcomes reported at 6, 12, and 24 months Total therapy hours: NR Manualized: NR Staff Training: NR Fidelity Checks: NR		
	Theory/Model: NR Program Type: Standard rehabilitation Setting: 4 general hospitals and 2 university teaching hospitals Delivery: N/A			

Appendix E. Table 2. Intervention Characteristics

Hashimoto, 2006¹⁰ Moderate to Severe TBI comprehensive treatment of varying intensities	Comprehensive Day Treatment program	Description: Group sessions focusing on enhancing individual's quality of life by teaching useful and effective behaviors and by redesigning patient's environment to help achieve goals. Coordination: All staff members Disciplines: Physical, social work, psychology, speech, vocational, "gymnastics," occupational, welfare Components: Therapy hours/week: 4 sessions/day for total of 4hrs/day for 6 months Duration: 6 months Total therapy hours: NR Manualized: NR Staff Training: NR Fidelity Checks: NR
	Theory/Model: Positivist-behavioral Program Type: Comprehensive Setting: Rehabilitation hospital Delivery: Group	Description: Group sessions focusing on enhancing individual's quality of life by teaching useful and effective behaviors and by redesigning patient's environment to help achieve goals. Coordination: All staff members Disciplines: Physical, social work, psychology, speech, vocational, "gymnastics," occupational, welfare Components: N/A Therapy hours/week: 4 sessions for total of 2 hrs/day, twice weekly Duration: 4 months Total therapy hours: NR Manualized: NR Staff Training: NR Fidelity Checks: NR

Appendix E. Table 2. Intervention Characteristics

Comprehensive Day Treatment program	Description: Group sessions focusing on enhancing individual's quality of life by teaching useful and effective behaviors and by redesigning patient's environment to help achieve goals.
Theory/Model: Positivist-behavioral	Coordination: All staff members
Program Type: Comprehensive	Disciplines: Physical, social work, psychology, speech, vocational, "gymnastics," occupational, welfare
Setting: Rehabilitation hospital	Components: Giving information, mentoring, goal-setting, reassurance, modeling problem-solving, referral to community resources, triaging to regional or tertiary center if local resources unavailable
Delivery: Group	Therapy hours/week: 4 sessions for total of 2 hrs/day, twice weekly
	Duration: 3 months
	Total therapy hours: NR
	Manualized: NR Staff Training: NR Fidelity Checks: NR
Comprehensive Day Treatment program	Description: Group sessions focusing on enhancing individual's quality of life by teaching useful and effective behaviors and by redesigning patient's environment to help achieve goals.
Theory/Model: Positivist-behavioral	Coordination: All staff members
Program Type: Comprehensive	Disciplines: Physical, social work, psychology, speech, vocational, "gymnastics," occupational, welfare
Setting: Rehabilitation hospital	Components: N/A
Delivery: Group	Therapy hours/week: 4 sessions for total of 2 hrs/day, twice weekly
	Duration: 4 months
	Total therapy hours: NR
	Manualized: NR Staff Training: NR Fidelity Checks: NR

Appendix E. Table 2. Intervention Characteristics

Ponsford, 2006⁵ Postacute moderate to severe TBI	Community-based therapy programme (CT)	Description: Access and conduct therapy in the home, workplace or community setting with active involvement of TBI individual, relatives and other s.
	Theory/Model: NR Program Type: Community-based group therapy Setting: Epworth Rehabilitation Programme (Australia) Delivery: Tailored to individual	Coordination: NR Disciplines: several disciplines; referrals made to local services; a significant number of patients do attend regular physiotherapy sessions at the rehabilitation center.. Components: Identification of important roles, goal setting, assessment of strengths and weaknesses, impairments and disabilities to be overcome to achieve goals. Therapies delivered in relevant setting. Therapy hours/week: NR, but most patients seen by a given therapist once a week or less Duration: NR Total therapy hours: NR. Manualized: NR Staff Training: NR Fidelity Checks: NR
	Hospital-based outpatient rehabilitation (historical)	Description: Group social communication skills training to improve pragmatic language skills, social behaviors and cognitive abilities.
	Theory/Model: NR Program Type: Hospital-based outpatient Setting: Epworth Rehabilitation Programme (Australia) Delivery: Tailored to individual	Coordination: NR Disciplines: NR Components: domain specific therapies and group sessions, visits to home, work, shopping, domestic activities. Therapy hours/week: NR Duration: NR Total therapy hours: NR Manualized: NR Staff Training: NR Fidelity Checks: NR

Appendix E. Table 2. Intervention Characteristics

Powell, 2002¹⁴ Chronic Severe TBI	Outreach	Description: a goal planning framework for delivering rehabilitation through individualized retraining delivered through community –based services.
	Theory/Model: NR Program Type: Multidisciplinary Outreach Setting: Homes or community settings – organized through Homerton Hospital (London) Delivery: Tailored to individual	Coordination: led by a clinical NP Disciplines: OT, PT, S:P, psych, SW Components:: Individual sessions, 2/week Therapy hours/week: 2-6 hours/week Duration: 6-12 weeks for goal setting/assessment; After initial assessment period, individuals seen for 27.3(sd=19.1) weeks for treatment Total therapy hours: NR Manualized: NR Staff Training: NR Fidelity Checks: NR
	Information	Description: One home visit by therapist who gave patient specially collated booklet listing resources and highlighting those relevant to patient's needs.
	Theory/Model: NR Program Type: Information Setting: Home - organized through Homerton Hospital (London) Delivery: Home visit & Standard booklet	Coordination: NR Disciplines: team therapist Components: Individual session, education Therapy hours/week: 0 Duration: 1 visit Total therapy hours: 1 Manualized: NR Staff Training: NR Fidelity Checks: NR

Appendix E. Table 2. Intervention Characteristics

Prigatano, 1984⁹ Chronic Severe Closed Head Injury Patients	Neuropsychological Rehabilitation Program (NRP) Theory/Model: Milieu based programs (Ben-Yishay 1982, Rosenbaum et al., 1978) Program Type: Hospital-based outpatient Setting: Presbyterian Hospital (Oklahoma City, US) Delivery: Peer groups progress through treatments	Description: Intensive, coordinated treatment emphasizing awareness and acceptance of impairments; cognitive retraining of select residual deficits and the development of compensatory skills. Coordination: NR Disciplines: NP, SLP, OT, PT, psychologist Components: Small group and individual sessions Therapy hours/week: 24 Duration: 6 mo. Total therapy hours: 576 Manualized: Yes Staff Training: NR Fidelity Checks: NR
<hr/> Untreated <hr/>		
Prigatano, 1994⁷ Chronic Moderate to Severe TBI with adequate potential to return to work	Neuropsychological Rehabilitation Program (NRP) Theory/Model: Intensive holistic cognitive rehabilitation/milieu program (Ben-Yishay et al., 1985) Neuropsychological rehabilitation (Ben-Yishay, et al., 1987) Program Type: Work Re-entry program Setting: Adult Day Hospital for Neurological Rehabilitation, Saint Joseph's Medical Center (Phoenix, AZ) Delivery: Peer groups progress through treatment	Description: A series of interdisciplinary therapies embedded in a milieu program that emphasizes a holistic approach. Teaching patients to be part of a small community encouraging cooperation and responsibility. Simulated natural setting. Individual learns along with others. TBI patients who underwent a specialty rehabilitation program; after 6-8 weeks of therapy, patients were integrated into 15-20 hours of work per week Coordination: NR Disciplines: PT, OT, SLP, cognitive therapy Components: individual therapies depending upon needs, individual psychotherapy, daily group psychotherapy, 'simulated' community interaction, protected work trial. Therapy hours/week: 24 Duration: 6 mo. Total therapy hours: approximately 576 Manualized: No Staff Training: NR Fidelity Checks: NR

Appendix E. Table 2. Intervention Characteristics

Untreated (historical)		
Rattok, 1992^s Cognitive remediation	Treatment 1 - Balanced	<p>Description: Balanced package that included training to alleviate attentional disorders, individualized cognitive remediation, small-group interpersonal communication exercises, therapeutic community activities, and personal counseling functions. Remediative cognitive training included.</p> <p>Coordination: NR</p> <p>Disciplines: NR</p> <p>Components: Individual and small-group counseling</p> <p>Therapy hours/week: 5hr/day, 4 days/week</p> <p>Duration: 20 weeks</p> <p>Total therapy hours: 200</p> <p>Manualized: NR Staff Training: NR Fidelity Checks: NR</p>
	Treatment 2 - Interpersonal	<p>Description: Training in attention, community activities, and personal counseling; no individualized counseling; emphasis on small-group interpersonal exercises</p> <p>Coordination: NR</p> <p>Disciplines: NR</p> <p>Components: Group work</p> <p>Therapy hours/week: 5hr/day, 4 days/week</p> <p>Duration: 20 weeks</p> <p>Total therapy hours: 200</p> <p>Manualized: NR Staff Training: NR Fidelity Checks: NR</p>

Appendix E. Table 2. Intervention Characteristics

Salazar, 2000³ Moderate to Severe Closed head injury among active duty military personnel	Inpatient Cognitive Rehabilitation Theory/Model: Milieu-oriented approach modified to fit military framework (Prigatano 1994 Prigatano 1989); intergrated work therapy (Ben-Yishay 1987, Burke 1988) Setting: minimum care hospital ward, Walter Reed Army Medical Center (Washington, DC) Delivery: Peer groups progress through treatmen	Description: In a military milieu, physical fitness training and group and individual cognitive, speech, occupational, and coping skills therapies conducted with integrated work therapy coordinated to simulate patient's previous work or military specialty Coordination: Psychiatrist Disciplines: Neuropsychology, occupational therapy, speech pathology, physical therapy, neurological and psychiatric consultation Components: Group and individual Therapy hours/week: NR Duration: 6 wks. Total therapy hours: NR Manualized: Yes Staff Training: NR Fidelity Checks: Intermittent reviews and continuing education
	Home rehabilitation Theory/Model: NR Program Type: Home-based postacute rehabilitation Setting: Home Delivery: Visits and phone calls from psychiatric nurse.	Description: Patients received TBI education and individual counseling from a psychiatric nurse and were given educational materials and recommended strategies for enhancing cognitive and organizational skills. included Disciplines: psychiatric nurse Components: Trained to in various home number and card games; encouragement to read and watch news programs, resumed daily physical exercise at their own pace. Therapy hours/week: .5 h/wk Duration weeds: 8 weeks Total therapy hours: NR Manualized: Yes Staff Training: NR Fidelity Checks: NR

Appendix E. Table 2. Intervention Characteristics

Sarajuuri, 2005^b Chronic Moderate to Severe TBI	INSURE Program	<p>Description: Postacute, interdisciplinary, 6-week, inpatient neuropsychologic rehabilitation and psychotherapy. Therapeutic alliance is emphasized. Compensatory techniques,</p> <p>Coordination: NR</p> <p>Disciplines: NP, neurologist, rehabilitation nurse, SW, SPL, OT, PT</p> <p>Components: Cognitive group – 2 session/wk, pragmatic group – 1 session/wk, pictures of self group – 1 session/ wk, quality of life group – 1 session/ wk, sport, relaxation, and jogging group – 1 session/ wk; 2-day seminar with participation from family, employers, public health professionals to plan remaining 2 wks of program; supported and individually tailored vocational interventions.</p> <p>Therapy hours/week: 37.5</p> <p>Duration weeks: 6 weeks</p> <p>Total therapy hours: 225</p> <p>Manualized: Yes Staff Training: NR Fidelity Checks: NR</p>
	<p>Theory/Model: Neuropsychologic rehabilitation and psychotherapy (Ben-Yishay 1987 ; Ben-Yishay 1985 Christensen 1992, Prigatano 1986)</p> <p>Program Type: Residential Neuropsychologic rehabilitation</p> <p>Setting: Kapyla Rehabilitation Centre (Helsinki, Finland)</p> <p>Delivery: Peer groups progress through treatment</p>	
	Conventional Rehabilitation	<p>Description: Conventional clinical care and rehabilitation in local healthcare system. Rehabilitation services individually tailored and delivered in an unstructured and nonsystematic way.</p> <p>Coordination: NR</p> <p>Disciplines: Such as PR, PR SLP, NP and psychotherapy</p> <p>Components: NR</p> <p>Therapy hours/week: NR</p> <p>Duration: NR</p> <p>Program total therapy hours: NR</p> <p>Manualized: No Staff Training: No Fidelity Checks: No</p>
	<p>Theory/Model: NR</p> <p>Program Type: As referred by physician</p> <p>Setting: Recruited from Department of Neurosurgery, Helsinki University Central Hospital, Level 1 Trauma Center</p> <p>Delivery: As referred by physician</p>	

Appendix E. Table 2. Intervention Characteristics

Semlyen, 1998¹⁶ Postacute Severe TBI	Multidisciplinary rehabilitation Theory/Model: NR Program Type: Residential Neuropsychologic rehabilitation Setting: Hunters Moor Regional Rehabilitation Centre (Newcastle upon Tyne, UK) Delivery: Coordinated, multidisciplinary rehabilitation delivered individually	Description: Coordinated multidisciplinary approach that could include Inpatient, outpatient or home-based services delivered by multidisciplinary team with TBI specialization and coordinated patient goal setting with patient, team, and family members. Weekly review of goals. Coordination: NR Disciplines: nursing, PT, SLP, OT, clinical psychology, rehabilitation medicine, counseling, social work Components: individualized, daily Therapy hours/week: NR Duration: 201.0±144.12 (mean days±SD); Total therapy hours: NR Manualized: NR Staff Training: NR Fidelity Checks: NR
	Single discipline approach Theory/Model: NR Program Type: variable Setting: settings other than Hunters Moor Regional Rehabilitation Centre (Newcastle upon Tyne, UK) Delivery: variable, but independent for each Individual	Description: Less coordinated, single discipline approaches including inpatient and outpatient rehabilitation and could be only physiotherapy delivered for 1 hour once a week or several therapies providing input several times a week. Coordination: NR Disciplines: NR Components: variable Total therapy hours/week: NR Program Duration: 111.80±175.17 (mean days±SD) Total therapy hours: NR Manualized: NR Staff Training: NR Fidelity Checks: NR

Appendix E. Table 2. Intervention Characteristics

Thomas, 2004¹⁵ Adjustment to Acquired Brain Injury	Potential Unlimited Program (PUP) Theory/Model: Simpson, 1996; Understanding, Re-integrating identity, acceptance, restructuring Program Type: Outward Bound Setting: Community, Outward Bound course (Australia), patient home Delivery: Mixed	Description: Three stage program consisting of 1)Group fundraising, 2)9-day Outward Bound “Discovery” course adapted to accommodate patients’ needs, 3)Follow-up group work to transfer insights from program to key areas of psychosocial functioning Coordination: NR Disciplines: NR Components: Goal setting, group work, physical activities Therapy hours/week: Stage 1 = NR, Stage 2= 9 days, Stage 3 = 2 hours every other week for 3-4 months Duration: NR Total therapy hours: NR Manualized: Outward Bound portion (Stage 2) Staff Training: NR Fidelity Checks: NR
	Control Theory/Model: NR Program Type: NR Setting: NR Delivery: N/A	Description: Matched patients who had expressed initial interest in the PUP but were unable to participate Coordination: NR Disciplines: NR Components: NR Therapy hours/week: NR Duration: Assessments taken at same time points as PUP group Total therapy hours: NR Manualized: NR Staff Training: NR Fidelity Checks: NR

Appendix E. Table 2. Intervention Characteristics

Vanderploeg, 2008² Postacute Moderate to Severe TBI in veterans or active duty military personnel]	Cognitive didactic treatments inpatient TBI rehabilitation	Description: Emphasized explicit learning in an environment permitting and encouraging errors to assist clients to develop cognitive self-awareness. Targeting specific cognitive processes. Targeted 4 cognitive domains (attention, memory, executive function, and pragmatic communication) using trial-and-error learning approach to address patient self-awareness. Directly rehabilitating the cognitive deficits that underlie most functional TBI deficits to result in a generalized functional improvement.
	Theory/Model: Cognitive-didactic treatments (Sohlberg & Mateer 1986, 1989, 2001) Program Type: Residential postacute rehabilitation center Setting: Four VA inpatient postacute rehabilitation centers Delivery: Individual in person	Coordination: Psychiatrist Disciplines: Rehabilitation nurses, PT, PR, rehabilitation counseling, patient and family education, psychologic or SW support services, Occupational therapy, physical therapy, speech/cognitive/language therapy, neuropsychology Components: 7.5-15 hrs/wk cognitive didactic treatment integrated into essential CARF standard of care interdisciplinary rehabilitation. Memory notebooks. Therapy hours/week: 21.5-30 hrs/wk Duration: 32.2(±12.2) days Total therapy hours: NR; continued until clinically judged ready for discharge or 60 days Manualized: No Staff Training: Yes Fidelity Checks: Yes
	Functional-experiential treatments within inpatient TBI rehabilitation	Description: Real life performance situations and common tasks to remediate or compensate for functional deficits Learning-by-doing functional daily activities using an errorless treatment strategy incorporating therapist direction and structure to complete components of gradually more complex tasks; did not entail explicit awareness or learning, but rather emphasized motor and other forms of implicit learning.
	Theory/Model: Functional treatment concepts (Giles 1993, 1999, 2006; Hartley 1995) Program Type: Residential postacute rehabilitation center Setting: Four VA inpatient acute rehabilitation centers Delivery: Groups in natural settings	Coordination: Psychiatrist Disciplines: Occupational therapy, physical therapy, speech/cognitive/language therapy, neuropsychology Rehab Goals: To use real-life performance situations and common tasks to remediate or compensate for functional deficits Components: 7.5-15 hrs/wk functional-experimental treatment integrated into essential CARF standard of care interdisciplinary rehabilitation. Memory notebooks. Therapy hours/week: 21.5-30 hrs/wk Duration: 33.3(±13.6) mean (std dev) days Total therapy hours: NR; continued until clinically judged for discharge or until 60 days Manualized: No Staff Training: Yes Fidelity Checks: Yes

Appendix E. Table 2. Intervention Characteristics

Willer, 1999¹² Postacute severe brain injury with multiple disabilities	Community-based residential rehabilitation Theory/Model: Cognitive rehabilitation and community readaptation (Fryer 1987) Program Type: Residential postacute rehabilitation program Setting: homelike residential (Canada) Delivery: Individuals	Description: TBI subjects who received postacute, community and residential-based rehabilitation Coordination: NP Disciplines: MD, PT, OT, SPL, paraprofessionals Components: NR Therapy hours/week: NR Duration: ≥ 1 year (up to 3 years) Total therapy hours: NR Manualized: No Staff Training: Yes Fidelity Checks: No
	Home-based rehabilitation services Theory/Model: NA Program Type: varies Setting: Home and outpatient services Delivery: Individuals	Description: A highly variable range of home-based or outpatient services. Coordination: NR Disciplines: occupational and physical therapists, neuropsychology, case management , and nursing services Components: NR Total therapy hours/week: NR Program Duration: ≥ 1 year (up to 3) Total therapy hours: NR Manualized: No Staff Training: Yes Fidelity Checks: No

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